<u>claims</u>

 A method for detecting a target nucleic acid molecule, said method comprises:

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- a) preparing a cell lysate comprising lysing a cell in a biological sample in a lysis buffer to release the target nucleic acid molecule from the cell;
- b) incubating the cell lysate from step a), without nucleic acid purification, with a nucleic acid probe immobilized on a solid substrate under conditions that allow hybridization between the target nucleic acid molecule and the probe, wherein the nucleic acid probe comprises a sequence complementary to the target nucleic acid molecule;
- c) assessing hybridization between the target nucleic acid molecule and the probe to determine the presence, absence and/or amount of the target nucleic acid molecule.
- 2. The method of claim 1, wherein the cell is lysed in the lysis buffer by a physical method.
 - 3. The method of claim 2, wherein the physical method is selected from the group consisting of grinding, ultrasonic lysing, lysing with high temperature, and freezing.
- 4. The method of claim 1, wherein the cell is lysed in the lysis buffer by a chemical method.
 - 5. The method of claim 4, wherein the chemical method is lysing with a protein denaturant or a detergent.
 - 6. The method of claim 1, wherein the cell is lysed in the lysis buffer by a biological method.
- 7. The method of claim 6, wherein the biological method is lysing with a proteinase or a lysozyme.
 - 8. The method of claim 1, wherein the cell is lysed by any combination of a physical, a chemical, and a biological method.

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9. The method of claim 1, wherein the cell lysate is incubated with the probe immobilized on the substrate in the lysis buffer for hybridization.

- 10. The method of claim 1, wherein an agent that aids for hybridization is added to the cell lysate before the cell lysate is incubated with the probe.
- 11. The method of claim 10, wherein the agent is selected from the group consisting of NaCl, citrate sodium, and SDS.

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- 12. The method of claim 1, wherein the biological sample is a sample selected from the group consisting of a non-virus biological organism, a biological tissue, a eukaryotic cell, and a prokaryotic cell.
- 13. The method of claim 1, wherein the target nucleic acid molecule is selected from the group consisting of a genomic DNA, a plasmid, a mitochondria DNA, a chloroplast DNA, a messenger RNA, a ribosomal RNA, and a small nuclear RNA.
 - 14. The method of claim 1, wherein the solid substrate comprises a material selected from the group consisting of a nylon film, a pyroxylin film, a silicon, a glass, a ceramic, a metal, a plastic, and a combination thereof.
 - 15. The method of claim 1, wherein the solid substrate comprises a plurality of nucleic acid probes, and wherein the plurality of the nucleic acid probes are immobilized on the solid substrate to form an array.
- 16. The method of claim 15, wherein the plurality of the nucleic acid probes have different nucleotide sequences.
 - 17. The method of claim 16, wherein the number of different probes is from about 2 to about 100,000.
 - 18. The method of claim 15, wherein the area of the array is from about 0.01 mm² to about 100 cm².
- 25 19. The method of claim 15, wherein the array is selected from the group consisting of a two-dimensional array, a three-dimensional array, and a four-dimensional array.

20. The method of claim 1, wherein the nucleic acid probe immobilized on the solid substrate comprises a single-stranded oligonucleotide or a double-stranded PCR product.

21. The method of claim 1, wherein the cell lysate comprises an agent selected from the group consisting of a detergent, a protein denaturant, a buffer, a nuclease inhibitor, a salt, and a combination thereof.

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22. The method of claim 1, wherein the hybridization between the target nucleic acid molecule and the nucleic acid probe is assessed by determining binding of a reporter to the target nucleic acid molecule, wherein the reporter comprises a detectable marker selected from the group consisting of a fluorescein, an isotope, a biotin, a digoxin, a gold colloid, a magnetic bead, a electrochemical label, and a chemiluminescent label.